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**(54) RECORDING SHEETS FOR INK JET PRINTING**

AUFZEICHNUNGSSCHICHT FÜR TINTENSTRAHLDRUCK

FEUILLE D'IMPRESSION POUR IMPRESSION A JETS D'ENCRE

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**EP 0 705 172 B1**

**Description**Field of Invention

5 This invention relates to ink jet recording sheets for use in an ink jet recording process, particularly it relates to ink receiving sheets where images recorded thereon can be observed by both reflected and transmitted light. Ink jet receiving materials used at the present time have a particular need for improvement in physical and handling properties, particularly in waterfastness and light stability as well as for improved image quality. A preferred embodiment of this invention is therefore directed towards ink jet recording materials with improved handling and performance characteristics, in particular ink receiving materials where the images recorded thereon are resistant to rubbing on the surface or to damage by other physical means, remain intact in contact to water and do not fade when exposed to light even under adverse conditions. The present invention provides a solution towards these problems.

10 Ink jet printing systems generally are of two types: continuous stream and drop-on-demand. In continuous stream ink jet systems, ink is emitted in a continuous stream under pressure through an orifice or nozzle. The stream is perturbed, causing it to break up into droplets at a fixed distance from the orifice. At the break-up point, the droplets are charged in accordance with digital data signals and passed through an electric static field which adjusts the trajectory of each droplet in order to direct it to a gutter for recirculation or a specific location on a recording medium. In drop-on-demand systems, a droplet is expelled from an orifice to a position on a recording medium in accordance with digital data signals. A droplet is not formed or expelled unless it is to be placed on the recording medium.

20 Although the main effort in this invention is directed towards the more demanding continuous stream system it is not meant to be restricted to either of the two methods.

Background Art

25 The following requirements describe some of the major features of a recording material used in ink jet printing:

1. Sufficient ink absorbing capacity and ink receptivity of the receiving layer to prevent the ink from streaking and from running down during printing, even under conditions where several droplets are deposited in a rapid sequence onto the same spot.
- 30 2. Fast drying of the layer surface after printing of the image leading to prints free from tackiness.
3. Excellent colour rendition, no change of the hue of the picture with time.
- 35 4. Surface with high gloss.
5. In the case of transparencies, clear, transparent, scatter free receiving layers.
- 40 6. Resistance of the image surface of the image to rubbing.
7. Excellent waterfastness of the produced images.
8. Excellent light fastness of the printed images.
- 45 9. Excellent archival stability.
10. Excellent physical and handling properties.

50 The particular problem of waterfastness has in the past been addressed by a wide variety of techniques. Thus solutions to the problem have been proposed for by specific formulations of the inks or alternatively in many cases by specific modifications of the receiving layers. The two approaches have in many cases been combined.

One attempt to improve waterfastness has been the use of reactive dyes. So for instance in US 4,443,223 (Kissling et al.), US 5,230,733 (Pawlowski et al.), JSDC (1993), 109,147 (S.O. Aston et al.) and references cited therein. Although some improvement has been achieved by this technique no satisfactory results can in general be obtained due to the fact that the conditions which are possible in practice in a printing environment are less than optimal and do in general not suffice to achieve reaction of these dyes with given binders. Inks based on colloidal dye dispersions and polymers in inks have been proposed so for instance in US 5,100,471 (Wink et al.), 5,017,644 (Fuller et al.), 4,990,186 (Jones et al.), US 4,597,794 (Kasha et al.), US 4,210,566 (Murrey), US 4,136,076 (Dennison et al.), US 5,224,987

(Matrick et al.), US 5,180,425 (Matrick et al.), and US 4,246,154 (Yao et al.). Inks based on colloidal dyes as well as hot melt inks, although yielding images with good waterfastness and good light stability, do in many cases lead to images which are not transparent and therefore less suited to be used for projections.

Often involved modifications of the inks have the tendency to give inks liable to show precipitates upon prolonged storage. Such precipitates subsequently tend to clog the nozzles of ink jet printer.

The major attempt to achieve waterfastness in receiving layers has been via the use of polymers, particular cationic polymers in conjunction with inks containing acidic dyes.

US 4,877,680 describes cationic polymers together with neutral binders. Cationically modified polyvinyl alcohol has been described in US 4,783,376. US 4,575,465 claims quaternised polyvinyl pyridine to achieve waterfastness. US 4,554,181 describes the use of a combination of cationic polymers and polyvalent metal salts since only such combinations and not the single elements tend to provide the sought for properties.

Although good waterfastness can in general be obtained with a wide variety of cationic polymers they tend to show a severe drawback in that they impair the light fastness of the printed images.

The introduction of inorganic pigments, fillers, minerals, metal salts and metal oxides have been proposed. US 4,116,910 (Rudolphy et al.) propose the use of derivatives of metals of Group II of the periodic table together with natural resin. JP 6025,7285 (Nakatsugawa et al.) claims an improvement of light stability by addition of transition metal oxides which are used as ultra-fine particles. Waterfastness can preferentially be achieved by addition of metal oxides together with cationic pigments or polymers to the receiving layers. US 5,104,730 (Misuda et al.) and US 4,879,166 (Misuda et al.) describe porous recording sheets where the porous layer is mainly made of pseudo boehmite, a colloidal aluminium oxide hydroxide. Although in general satisfactory waterfastness can be achieved the layers obtained by this method are slightly opaque and show severe tendency to become brittle with time and on exposure to light.

All the above mentioned solutions fulfil only partly the requirements of image receiving layers for modern ink jet printing. In many cases these solutions lead moreover only to material suitable for quite restricted applications. Improvements incorporated into ink receiving layers, widely applicable to modern ink jet printing technology, are therefore the scope of this invention.

#### Disclosure of the Invention

Accordingly, an object of the present invention is to provide image receiving media for use in ink jet recording which are particularly excellent in water resistance.

Another object of the invention is to provide ink-jet recording media which achieve waterfastness without affecting the stability of the obtained images against the detrimental effect of light.

A further objective is to obtain ink-jet recording layers with excellent surface properties showing high resistance to physical damage like for instance scratching, resistance to cracking and moist rubbing on the surface.

Another objective of the invention is to obtain ink-jet recording media which satisfy in sharpness and surface lustre of the recorded image and are free from stickiness of the surface even under highly humid conditions.

A further objective is to provide ink-jet recording media, to be used on ink jet printers of the continuous stream type. A further object is to provide ink-jet media which allow the possibility to print images intended to match those on silver halide photographic material.

It is furthermore the objective of this invention to provide clear, scatter free recorded ink-jet images on transparent base material intended to be projected.

An additional objective of this invention is to obtain an ink-jet recording medium for office desk top publishing of colour graphics which has improved lightfastness.

This invention proposes to achieve above objectives by providing an ink-jet recording sheet wherein said sheet consists of a support, opaque or transparent, onto which has been coated a receiving layer or layers comprising a binder or a mixture of different binders, fillers, natural or synthetic polymers and wherein are imbedded or coated at least one water-soluble salt of Yttrium, Lanthanum, Cerium, Neodymium or Ytterbium.

These salts are hereinafter referred to as the selected metal salts of Group IIIB.

The preferred salts are salts of mineral acids and in particular nitric acid.

To these layers can be added a wide variety of additional elements to further improve the pictorial or/and physical properties of the images obtained when printed on an ink jet printer.

#### Best Mode of Carrying Out the Invention

The present invention will now be described in detail. The ink jet recording sheets according to this invention specifically relate to layers wherein are imbedded or coated with the selected water soluble salts of Group IIIB metals.

Preferably Lanthanum nitrate is present in the layer.

It is to be understood that the phrase 'selected metal salts of Group IIIB or complexes thereof' specifically excludes

oxides of such metals. The metal salts used in this invention can be present as halides, salts of most oxo acids, sulphates, nitrates, perchlorates, bromates but also as carbonates, phosphates or hydroxides. Also salts of organic acids can be used.

Often the salts are present as dissociated hydrated species and are in general used as such. The salts of the invention can also be used as mixtures made up of single species. There is no limitation as far as the ratios of the mixtures are concerned.

The salts used in the invention can be used in form of their double salts containing besides the selected metals of Group IIb the elements Ca, Mg, Ba, Na, K or the like. Double salts can be in form of, for instance, sulphates, nitrates, phosphates or in other forms known to those skilled in the art. The salts of the selected compounds when readily water soluble are introduced into the receiving sheets as aqueous solutions. In many instances the selected metal derivatives are only sparingly soluble in water and have to be applied in colloidal form or in form of fine dispersions.

The selected salts of Group IIb elements coated directly on the substrate or incorporated into the ink receiving layers of the proposed ink-jet recording material are added in an amount of 0.05 - 3.0g/m<sup>2</sup>, preferentially in an amount of 0.1 to 0.9g/m<sup>2</sup>. In the embodiment where the selected salts of Group IIb elements are coated directly on the substrate, they are preferably applied as 3 - 5% aqueous solutions on the surface of the substrate. After evaporation of the aqueous solution the salts essentially are absorbed into the substrate surface.

In this embodiment a preferred substrate includes a base paper sheet coated with a silica and polyvinyl alcohol matrix prior to application of the coating solution.

The use of the selected salts of the group IIb elements perform most efficiently when they are imbedded into layers or coated onto substrates which have the ability to rapidly absorb aqueous inks. The absorbing power of the layer is to a great extent a function of the materials used but likewise of the physical properties of the layers and the substrate. The compounds that make up the imbedding matrix include in general water soluble film forming polymers.

These film forming water soluble polymers may include, for example, natural polymers or modified products thereof such as albumin, gelatine, casein, starch, gum arabic, sodium alginate, hydroxyethyl cellulose, carboxymethyl cellulose,  $\alpha$ -,  $\beta$ - or  $\gamma$ -cyclodextrine and the like; polyvinyl alcohol; complete or partially saponified products of copolymers of vinyl acetate and other monomers; homopolymers or copolymers with other monomers of unsaturated carboxylic acids such as (meth) acrylic acid, maleic acid, crotonic acid and the like; homopolymers or copolymers with other vinyl monomers of sulfonated vinyl monomers such as vinylsulfonic acid, sulfonated styrene and the like; homopolymers or copolymers with other vinyl monomers of (meth) acrylamide; homopolymers or copolymers with other monomers of ethylene oxide; polyurethanes, polyamides having such groups as mentioned above; polyethyleneimine, polyacrylamides, water soluble nylon type polymers, polyvinylpyrrolidone, polyester; and so on. All these can also be used in mixtures.

These polymers can be blended with non water soluble natural or synthetic high molecular compounds.

Suitable synthetic polymer materials can be chosen from among polyvinyl lactams, acrylamide polymers, polyvinyl alcohol and its derivatives, polyvinyl acetals, polymers of alkyl and sulfoalkyl acrylates and methacrylates, hydrolyzed polyvinyl acetates, polyamides, polyvinyl pyridines, acrylic acid polymers, maleic anhydride copolymers, polyalkylene oxides, methacrylamide copolymers, polyvinyl oxazolidinones, maleic acid copolymers, vinylamine copolymers, methacrylic acid copolymers, acryloyloxyalkylsulfonic acid copolymers, sulfoalkylacrylamide copolymers, polyalkyleneimine copolymers, polyamines, N,N-dialkylaminoalkyl acrylates, vinyl imidazole copolymers, vinyl sulphide copolymers, halogenated styrene polymers, amineacrylamide polymers, polypeptides and the like.

Non-water soluble polymers can also be used in some cases.

In the case where one of the water soluble polymers is gelatine the types of gelatine suitable for use in the present invention include all kinds of gelatine currently known, for instance acid pigskin or limed bone gelatine, acid or base hydrolysed gelatines, but also derivatised gelatines like for instance phthalated, acetylated or carbamoylated, or gelatine derivatives with trimellitic acid. The preferred gelatine is a gelatine with an isoelectric point between 7 and 9.5.

The polymers mentioned above having reactive groups or groups having the possibility to react with a crosslinking agent can be cross linked to form essentially non water soluble layers. Such crosslinking bonds may be either covalent or ionic. Thus crosslinking allows for the modification of the physical properties of the layers, like for instance in water absorbency of the layer, but also in resistance against physical damage.

Crosslinking agents suitable for this particular use are selected depending on the water soluble polymer used. They may include for example chromium salts (such as chrome alum or chromium acetate), aldehydes (such as formaldehyde, glyoxal or glutaraldehyde), N-methylol compounds (such as dimethylolurea or methylol-dimethylhydantoin), dioxane derivatives (such as 2,3-dihydroxydioxane), activated vinyl compounds (such as 1,3,5-trisacryloyl-hexahydro-s-triazine or bis(vinylsulfonyl)methyl ether), activated halogen compounds (such as 2,4-dichloro-6-hydroxy-s-triazine), amino or substituted-amino modified triazines, epoxides, carbamoyl-pyridinium compounds or mixtures of two or more of above mentioned crosslinking agents.

The layers and coatings can be modified by addition of fillers. Possible fillers of the kind are for instance clay, kaolin, talcum, Ca-, Mg- or Ba-carbonate, Ca- or Ba-sulphate, silica, titanium oxide, chalk, bentonite, zeolite, aluminium silicate, calcium silicate, silicium oxide, satin white, colloidal silicium oxide and the like. Likewise the possibility exists to

use organic inert particles such as polymer beads. This includes beads made from polyacrylates, polystyrene or different copolymers of acrylates and styrene. These fillers are selected according to the intended use of the printed image. Some of these compounds cannot be used if the printed image is to be used as a transparency. Alternatively they are of interest in cases where the printed image is to be used as a reflected image. Often the introduction of such filler causes a desired matte surface.

The ink-jet image recording elements of this invention comprise a support for the ink receiving layer. A wide variety of such supports are known and commonly employed in the art. They include, for example, those supports used in the manufacture of photographic clear films including cellulose esters such as cellulose triacetate, cellulose acetate propionate or cellulose acetate butyrate, polyesters such as poly(ethylene terephthalate), polyamides, polycarbonates, polyimides, polyolefins, poly(vinyl acetals), polyethers, polyvinyl chloride and polysulfonamides. Polyester film supports, and especially poly(ethylene terephthalate) are preferred because of their excellent dimensional stability characteristics.

Likewise the usual supports commonly used in manufacturing of opaque photographic material can be used according to the present invention. They include baryta paper, polyethylene-coated paper, polypropylene synthetic paper, voided polyester as for instance manufactured by ICI under the trade name of MELINEX as well as voided polypropylene polyester likewise manufactured by the same company. Preferred are clear polyester, acetate, voided polyester or resin coated paper. When such support material, in particular polyester, is used a subbing layer is advantageously added first to improve the bonding of the ink receiving layer to the support. Useful subbing compositions for this purpose are well known in the photographic art and include, for example, polymers of vinylidene chloride such as vinylidene chloride/acrylonitrile/acrylic acid terpolymers or vinylidene chloride/methyl acrylate/itaconic acid terpolymers. Also usable are plain paper, comprising a wide variety of sizings, cast-coated papers and aluminium foils.

In certain embodiments of the invention, a preferred substrate includes a base paper sheet coated with a silica and polyvinyl alcohol matrix. When such support material is used an aqueous coating of metal salts of the selected Group IIb elements may be coated directly on the substrate surface. The inclusion of a film forming polymer in this coating formulation is optional when the described substrate or a similar one is used. This embodiment provides a recording medium suitable for office desk top publishing of colour graphics and has improved lightfastness properties.

The ink-jet ink receiving layers or coatings according to this invention are in general coated from aqueous solutions containing binders, additives, pigments and the like as well as the metal salts of use in the present invention. It is in many cases necessary to add surfactants to those coating solutions allowing for smooth coating and evenness of the layers.

Examples of suitable surfactants are non-ionic surface active agents such as saponin (steroids), alkylene oxide derivatives (such as polyethylene glycol, polyethylene glycol/polypropylene glycol condensates, polyethylene glycol alkyl or alkylaryl ethers, polyethylene glycol esters, polyethylene glycol sorbitan esters, polyalkylene glycol alkylamine or amides or silicone/polyethylene oxide adducts), glycidol derivatives (such as alkenylsuccinic acid polyglycerides or alkylphenol polyglycerides), aliphatic esters of polyhydric alcohols, alkyl esters of sucrose, urethanes or ethers; a sulfuric acid ester group or a phosphoric acid ester group, such as triterpenoid type saponin, alkylcarboxylates, alkylsulfonates, alkylbenzenesulfonates, alkyl naphthalenesulfonates, alkyl sulfuric acid esters alkyl phosphoric acid esters, N-acyl-N-alkyltaurines, sulfosuccinates, sulfo-alkylpolyoxyethylene alkylphenyl ethers or polyoxyethylene alkyl-phosphates and cationic surface active agents such as alkylamine salts, aliphatic or aromatic quaternary ammonium salts (such as pyridinium or imidazolium salts) or phosphonium or sulfonium salt containing an aliphatic or heterocyclic ring. Equally suitable are fluorinated or perfluorinated derivatives of the above mentioned compounds.

Specific examples of these surface active agents are those described in, e.g. US Pat. Nos. 2,240,472, 1,831,766, 3,158,484, 3,210,191, 3,294,540 and 3,507,660, British Pat. Nos. 1,012,495, 1,022,878, 1,179,290 and 1,198,450, US Pat. Nos. 2,739,891, 2,823,123, 3,068,101, 3,415,649, 3,666,478 and 3,756,828, British Pat. No. 1,397,218, US Pat. No. 1,397,218, US Pat. Nos. 3,133,816, 3,441,413, 3,475,174, 3,545,974, 3,726,683, and 3,843,368, Belgium Pat. No. 731,126, British Pat. Nos. 1,138,514, 1,159,825 and 1,374,780, 2nd US Pat. Nos. 2,271,623, 2,288,226, 2,944,900, 3,253,919, 3,671,247, 3,772,021, 3,589,906 and 3,754,924.

Besides being necessary for coating purpose surfactants may have an influence on the quality of the generated images and may therefore be selected with this specific goal in mind. There is in general no limitation to the use of the types of surfactants used as long as they do not interfere with the metal salts used in the present invention and later with the printing inks used for the production of the image.

Typically the ink-jet ink receiving layers according to this invention have a thickness in the range of 0.5 to 30 micrometers, preferably in the range of 2.0 to 15 micrometers dry thickness.

The coating solutions or coating dispersions can be coated onto a support by any number of suitable procedures. Usual coating methods include immersion or dip coating roll coating, air knife coating, extrusion doctor blade coating, rod and/or blade metering, or by spraying. An ink receiving system can be built up by several layers.

These layers can be coated one after the other or simultaneously. It is likewise possible to coat a support on both sides with ink receiving layers. Alternatively the backside may be coated with auxiliary layers like for instance anticurl

layers or antistatic layers. The way however by which the claimed receptive layers are produced is not to be considered limiting for the present invention. In addition to the above mentioned elements ink receiving layers as claimed in this invention can contain additional additives aimed at improving appearance as well as performance of the produced imaging material. It can for instance be beneficial to add brightening agents to the layers of receiving sheets. There is in general no limitation as to the kind of brighteners used. Suitable brightening agents are for instance stilbenes, coumarines, triazines or oxazoles or others known in the art.

Light stability can in general be improved further by adding UV absorbers to the layers. Although UV absorbers are in general added to the topmost layer of the system there is no limitation as to where within the ink receiving element such light absorbing compounds are added. The amount of UV-absorber can vary from 200-2000mg/m<sup>2</sup>, preferably however from 400mg to 1000mg/m<sup>2</sup>. Suitable types of absorbers can be for example benztriazoles, benzophenones, derivatives of acrylonitrile, thiazolidone, oxazole and thiazole.

It is further known that images can be protected from degradation by the addition of light stabilisers and antioxidants. Examples of such compounds are among others sterically hindered phenols, sterically hindered amines, chromanols and the like. Above mentioned additives can, if water soluble, be added as aqueous solutions. In the case where these compounds are not water soluble the above mentioned additives can be incorporated in the ink receiving element by common techniques known in the art. The compound is typically dissolved in a solvent selected from organic solvents compatible with water, such as alcohols, glycols, ketones, esters, amides and the like. Alternatively the compounds can be added to the layer as fine dispersions, as oil emulsions, as cyclodextrine inclusion complex or loaded as fine dispersions on to latex particles. Ultrasound or milling can be used to dissolve or disperse marginally soluble additives.

Inks for ink jet printing are well known. These inks consist in essence of a liquid vehicle and dissolved or suspended therein a dye or pigment. The liquid vehicle of the inks employed for the printing according to the present invention consist in general of water or a mixture of water and a miscible organic component such as ethylene glycol, and higher molecular glycols, glycerine, dipropylene glycol, polyethylene glycol, amides, polyvinylpyrrolidone, N-methylpyrrolidone, cyclohexylpyrrolidone, carboxylic acids and esters, ethers, alcohols, organosulfoxides, sulfolane, dimethylformamide, dimethylsulfoxide, cellosolve, polyurethanes, acrylates and the like.

The non-water part of the printing ink generally serves as humectant, cosolvent, viscosity regulating agent, ink penetration additive, levelling agent or drying agent. The organic component has in most cases a boiling point which is higher than that of water. In addition aqueous inks may contain inorganic or organic salts to impart electrical conductivity. Examples of such salts include nitrate, chlorides, phosphates and the like and salts of low molecular, water soluble organic acids like acetates, oxalates and similar. The dyes and pigments suitable for the preparation of inks usable with the receiving sheets of this invention cover practically all classes of known colouring compounds. Dyes or pigments typically used for that purpose are described in EP 0,559,324 (Isganitis et al.).

Other additives present in usable inks are for instance surfactants, optical brighteners UV absorbers or light stabilisers, biocides and polymeric additives. This description of inks is for illustration only and not to be considered as limiting the invention.

Test sheets prepared according to the described Examples were printed in an ink jet printer. 1cm by 1cm uniform patches were printed in cyan, magenta, yellow and black to a density of about 2. After printing and drying under ambient conditions for 12 hours the densities of the individual patches were measured with an X-rite densitometer.

#### Waterfastness

The samples printed and dried as described above were placed in deionised water at 20°C for 1 minute. The samples were then removed from the water, allowed to drip dry and remeasured. The difference between the densitometer readings was reported as % loss of optical density and termed waterfastness.

#### Light Stability

Sample sheets printed and dried as described above were exposed in an Atlas Weather-Ometer with a 2500 W Xenon lamp under conditions analogue to those set for in ISO norm 10 977. The samples were exposed until a total illumination of 40kj (kJoule)/cm<sup>2</sup> was reached. The difference between the densitometer readings was reported as % loss of optical density and termed light stability.

#### Example 1

18g gelatine with an isoelectric point of over seven (Stoess type 70810) were dissolved in 360ml deionized water. To this solution were added 12g hydroxyethyl cellulose. (Tylose H20, obtained from Hoechst AG) and 1.0g of a surfactant (Olin 10G, obtained from Olin Corporation). This solution was divided into twelve equal portions and to each por-

tion was added the amount of metal-nitrate x H<sub>2</sub>O indicated in Table 1. This amount corresponds to 0.125 mMol nitrate-salt/g total binder. A control solution contained no salt. Immediately before coating 0.55g of a 3% solution of 2-(4-dimethyl-carbamoyl-pyridino)-ethane-sulfonate was added to each portion. These solutions were then coated onto a subbed polyester support using a barcoater. The final dry thickness of the layers were approximately 8μ. The sheets were printed on an IRIS ink jet printer model 3024 with standard Iris writing fluids. The printed sheets were then treated as described in the above testing procedures. The obtained results are reported in Table 1.

Table 1

| Metal-Salt x H <sub>2</sub> O                       | g/g Binder | Key            | Waterfastness Loss of Density in % of Initial Density |    |    |    |
|---|------------|----------------|---|----|----|----|
|   |            |                | C   | H  | Y  | K  |
| La(NO <sub>3</sub> ) <sub>3</sub> 6H <sub>2</sub> O | 0.054      | i              | <1  | 4  | 8  | 6  |
| Yb(NO <sub>3</sub> ) <sub>3</sub> 5H <sub>2</sub> O | 0.056      | i              | <1  | 3  | 11 | 5  |
| Ce(NO <sub>3</sub> ) <sub>3</sub> 6H <sub>2</sub> O | 0.054      | i              | <1  | 3  | 9  | 8  |
| Y(NO <sub>3</sub> ) <sub>3</sub> 5H <sub>2</sub> O  | 0.046      | i              | <1  | 4  | 7  | 9  |
| Mg(NO <sub>3</sub> ) <sub>2</sub> 6H <sub>2</sub> O | 0.032*     | c <sup>2</sup> | 19  | 17 | 36 | 27 |
| Ba(NO <sub>3</sub> ) <sub>2</sub>                   | 0.033*     | c              | 21  | 12 | 27 | 27 |
| Ca(NO <sub>3</sub> ) <sub>2</sub> 4H <sub>2</sub> O | 0.030*     | c              | 27  | 18 | 33 | 27 |
| Zn(NO <sub>3</sub> ) <sub>2</sub> 6H <sub>2</sub> O | 0.037*     | c              | 26  | 20 | 39 | 29 |
| Al(NO <sub>3</sub> ) <sub>3</sub> 9H <sub>2</sub> O | 0.047      | c              | 45  | 11 | 13 | 29 |
| None  | 0          | c              | 36  | 20 | 31 | 36 |
| KEY:  |            |                |   |    |    |    |
| i <sup>1</sup> : Invention                          |            |                |   |    |    |    |

c<sup>2</sup>: Comparison

\* Coatings with most of the comparative salts were cloudy and could not possibly be used for transparent ink receiving material.

From the results in Table 1 can be seen that excellent waterfastness can be achieved with recording media according to the present invention while appreciable dye bleeding occurred with salts according to the state of the art.

### Example 2

Ink jet receiving sheets were prepared and tested in an analogous way as described in Example 1. In two cases the metal salts were replaced by cationic polymeric mordants (US 4,575,465) as indicated in Tables 2.1 and 2.2. The obtained results are reported in Tables 2.1 and 2.2.



Table 2.1

| Metal-Salt x H <sub>2</sub> O                       | g/g Binder | Key            | Waterfastness Loss in Density in %<br>after min. in water |    |    |    |
|---|------------|----------------|---|----|----|----|
|   |            |                | C   | H  | Y  | K  |
| La(NO <sub>3</sub> ) <sub>3</sub> 6H <sub>2</sub> O | 0.054      | i              | <1  | 5  | 9  | 8  |
| Y(NO <sub>3</sub> ) <sub>3</sub> 5H <sub>2</sub> O  | 0.056      | i              | <1  | 3  | 10 | 8  |
| Ce(NO <sub>3</sub> ) <sub>3</sub> 6H <sub>2</sub> O | 0.054      | i              | <1  | 6  | 9  | 13 |
| Mg(NO <sub>3</sub> ) <sub>2</sub> 6H <sub>2</sub> O | 0.032      | c <sup>2</sup> | 18  | 13 | 31 | 23 |
| Mordant 1 <sup>3</sup>                              | 0.08       | c              | 12  | 30 | 33 | 27 |
| Mordant 2 <sup>3</sup>                              | 0.08       | c              | 5   | 12 | 21 | 11 |
| None  |            |                | 37  | 15 | 29 | 31 |
| KEY:<br>i <sup>1</sup> : Invention                  |            |                |   |    |    |    |

c<sup>2</sup>: Comparison  
<sup>3</sup>: US 4,575,465

Table 2.2

| Metal-Salt x H <sub>2</sub> O                       | g/g Binder | Key            | Light stability Loss in Density %<br>after 20 kj Atlas |    |    |    |
|---|------------|----------------|--|----|----|----|
|   |            |                | C  | H  | Y  | K  |
| La(NO <sub>3</sub> ) <sub>3</sub> 6H <sub>2</sub> O | 0.054      | i              | 1  | 13 | 30 | 39 |
| Y(NO <sub>3</sub> ) <sub>3</sub> 5H <sub>2</sub> O  | 0.056      | i              | 4  | 11 | 30 | 40 |
| Ce(NO <sub>3</sub> ) <sub>3</sub> 6H <sub>2</sub> O | 0.054      | i              | 0  | 13 | 27 | 39 |
| Mg(NO <sub>3</sub> ) <sub>2</sub> 6H <sub>2</sub> O | 0.032      | c <sup>2</sup> | 9  | 13 | 32 | 31 |
| Mordant 1 <sup>3</sup>                              | 0.08       | c              | 9  | 65 | 47 | 65 |
| Mordant 2 <sup>3</sup>                              | 0.08       | c              | 5  | 93 | 50 | 87 |
| None  |            |                | 9  | 14 | 39 | 37 |
| KEY:<br>i <sup>1</sup> : Invention                  |            |                |  |    |    |    |

c<sup>2</sup>: Comparison  
<sup>3</sup>: US 4,575,465

From the results given in Table 2.1 the efficacy of the salts claimed in this invention in improving the waterfastness of the dyes in printed images is evident. It can furthermore be seen from table 2.1 and 2.2 that the efficacy in improving waterfastness by cationic mordants according to the state of the art is considerably lower than with the salts according to this invention. What is however particularly evident is that no deterioration of light stability occurs in presence of these salts where however the stability against light is completely lost in presence of these mordants.

### Example 3

A coating mixture with a solid content of about 20% was prepared as follows, comprising:

|    |  |       |
|----|--|-------|
| 1. | Gelatine (Stoess type 69 426)                        | 2.4g  |
| 2. | Polyurethan (Daothan 1226 Hoechst, 40% aqueous sol.) | 3.0g  |
| 3. | Kaolin   | 10.0g |
| 4. | Olin 10G (Surfactant, Olin Corp.)                    | 0.1g  |
| 5. | Crosslinker (idem Example 1)                         | 0.05g |
| 6. | La(NO <sub>3</sub> ) <sub>3</sub> · H <sub>2</sub> O | 5.0g  |
| 7. | Water to 100g  |       |

This mixture was bar coated onto an unsized high quality paper in an amount of 1.2g/m<sup>2</sup> (sample A). A control (sample B) was prepared in an analogous way but without the addition of Lanthanum salt. Waterfastness and light stability were determined as described above. The results are shown in Table 3.1.

Table 3.1

| Sample | Waterfastness: % Loss (1 Minute Water) |    |    | Light Stability: % Loss (20 KJ Atlas) |   |    |    |    |
|--------|--|----|----|---------------------------------------|---|----|----|----|
|        | C                                      | M  | Y  | K                                     | C | M  | Y  | K  |
| A      | <1                                     | 5  | 41 | <1                                    | 3 | 56 | 27 | 27 |
| B      | 13                                     | 25 | 87 | 31                                    | 7 | 67 | 50 | 60 |

The same samples were prepared but coated onto heavy weight water colour paper. The results are shown in Table 3.2.

Table 3.2

| Sample | Waterfastness: % Loss (1 Minute Water) |    |    | Light Stability: % Loss (20 KJ Atlas) |   |    |    |    |
|--------|--|----|----|---------------------------------------|---|----|----|----|
|        | C                                      | M  | Y  | K                                     | C | M  | Y  | K  |
| A      | <1                                     | <1 | 14 | <1                                    | 3 | 35 | 33 | 33 |
| B      | 13                                     | 32 | 73 | 39                                    | 6 | 32 | 68 | 66 |

The results in Table 3.1 and 3.2 clearly show the effect of the lanthanum salt on diffusion of the dyes in water, also in the case where the claimed system is applied to plain paper. Light stability is in both cases improved when compared to the sample not containing lanthanum salt.

#### Example 4

In this example lanthanum nitrate coatings were applied directly to a substrate surface and were evaluated for lightfastness and tendency to fade.

Four samples A, B, C & D were prepared. The substrate used in all samples is a base paper sheet coated with a silica and polyvinyl alcohol matrix. Particular physical details of the support are as follows:

24lb base sheet (Lock Haven - alkaline, wood fiber matrix - 60/40 hardwood/softwood) 25% CaCO<sub>3</sub> filler (precipitated HO/LO); internally sized with ASA; surface sized with starch.

Physical properties of the base sheet include:

|  |              |
|--|--------------|
| Basis Weight                           | 10kg (24 lb) |
| Caliper (mils 0.001 inch)              | 4.0          |
| Moisture (percent)                     | 4.8±.5       |
| Sheffield Smoothness (Sheffield units) | 40           |
| Porosity (Gurley)                      | 60           |

The base sheet has a brightness (GE percent) of 90+ and opacity (percent) of 94.

Strength properties include: Stiffness (Gurley); 2mgf, Tear MD & CD both 50+ g and Mullen 2 bar (30+ psi.)

Silica coating: fumed silica 30 parts; precipitated silica 70 parts; polyvinyl alcohol 40 parts; dispersant and surfactants 3.1 parts. This coating is applied to the base sheet in the range of 5 - 8kg/m<sup>2</sup> (3-5 lbs/3,000 ft.)

The silica coating is applied to the base sheet in 2 applications using a rod coater. A lanthanum nitrate (water soluble salt of Group IIb) is applied to the surface of the silica coating in either a 4% (Sample B) or a 3% (Sample C) aqueous solution using a rod coater. Other Group IIb metals that may be used include, yttrium, cerium, neodymium, and ytterbium. The backside coating of the substrate consists of a 0.5% calcium stearate solution that is used as an anticurl agent and to reduce the coefficient of friction.

The support without any coating (Sample A), and coated with a quarternary amine dye fixative (Sample D), were used as controls. The amine dye fixative coating formulation includes a quaternised amine ester, lauryldimethylbenzylammoniumchloride, a polyamine salt aqueous solution and a silicone derivative.

#### Lightfastness Evaluation

The effect of lanthanum nitrate on ink jet printing ink colour was evaluated before and after lightfastness testing. Hewlett Packard 560 series ink jet printing ink colour was used in this example but any other commercially available colour inks are also suitable for use in the invention. Samples A, B, C and D were exposed to carbon arc light for periods of 1, 2 and 4 hours. Colour readings (L\*A\*B\*) were taken on exposed and unexposed areas. The results are shown in the Table 4.1 below.

Table 4.1

| EFFECTS OF TOP COATING ON COLOUR (L*A*B*) |         |       |        |        |
|---|---------|-------|--------|--------|
| CONDITION                                 | COLOUR  | L*    | A*     | B*     |
| A - CONTROL                               | BLACK   | 34.43 | -12.04 | -6.49  |
| B   | BLACK   | 40.70 | -8.31  | -8.43  |
| C   | BLACK   | 39.69 | -8.89  | -7.13  |
| D - CONTROL                               | BLACK   | 32.43 | -6.64  | -8.00  |
| A - CONTROL                               | CYAN    | 50.85 | -33.09 | -54.05 |
| B   | CYAN    | 50.05 | -30.65 | -53.91 |
| C   | CYAN    | 51.30 | -32.51 | -53.59 |
| D - CONTROL                               | CYAN    | 49.71 | -27.07 | -56.35 |
| A - CONTROL                               | YELLOW  | 89.78 | -1.76  | 104.30 |
| B   | YELLOW  | 89.31 | -0.53  | 100.81 |
| C   | YELLOW  | 89.66 | -1.70  | 102.51 |
| D - CONTROL                               | YELLOW  | 88.28 | -0.63  | 98.58  |
| A - CONTROL                               | MAGENTA | 48.88 | 60.84  | -53.20 |
| B   | MAGENTA | 47.56 | 57.56  | -54.58 |

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Table 4.1 (continued)

| EFFECTS OF TOP COATING ON COLOUR (L*A*B*)   |         |       |       |        |
|---|---------|-------|-------|--------|
| CONDITION   | COLOUR  | L*    | A*    | B*     |
| C   | MAGENTA | 47.59 | 59.24 | -54.94 |
| D - CONTROL   | MAGENTA | 50.67 | 59.80 | -47.09 |
| KEY:<br>Sample A - Control: no lanthanum nitrate<br>Sample B - 0651 g/m <sup>2</sup> (0.4 lb/3,000 sq.ft.) lanthanum nitrate<br>Sample C - 0.488 g/m <sup>2</sup> (0.3 lb/3,000 sq.ft.) lanthanum nitrate<br>Sample D - Control: quarternary amine dye fixative |         |       |       |        |

The tendency to fade of the coated samples above were also evaluated after exposing treated papers to 1, 2 and 4 hours of carbon arc light. The results are shown in Table 4.2 below.

Table 4.2

| THE EFFECTS OF TOP COATING ON FADE<br>SAMPLE A (CONTROL) |                  |                  |                  |
|--|------------------|------------------|------------------|
| COLOUR   | 1 HOUR FADE D.E. | 2 HOUR FADE D.E. | 4 HOUR FADE D.E. |
| BLACK  | 7.78             | 10.53            | 25.64            |
| CYAN   | 13.91            | 18.58            | 34.54            |
| YELLOW   | 3.10             | 3.71             | 6.39             |
| MAGENTA  | 18.98            | 30.54            | 46.80            |
| AVERAGE FADE - 18.29                                     |                  |                  |                  |
| SAMPLE B (4% La(NO <sub>3</sub> ) <sub>3</sub> )         |                  |                  |                  |
| COLOUR   | 1 HOUR FADE D.E. | 2 HOUR FADE D.E. | 4 HOUR FADE D.E. |
| BLACK  | 3.79             | 6.11             | 11.73            |
| CYAN   | 18.78            | 24.33            | 35.42            |
| YELLOW   | 2.20             | 2.70             | 3.72             |
| MAGENTA  | 20.13            | 29.58            | 44.97            |
| AVERAGE FADE - 16.95                                     |                  |                  |                  |
| SAMPLE C (3% La(NO <sub>3</sub> ) <sub>3</sub> )         |                  |                  |                  |
| COLOUR   | 1 HOUR FADE D.E. | 2 HOUR FADE D.E. | 4 HOUR FADE D.E. |
| BLACK  | 5.28             | 7.74             | 16.39            |
| CYAN   | 16.09            | 22.19            | 33.35            |
| YELLOW   | 3.0              | 3.86             | 5.78             |
| MAGENTA  | 20.03            | 30.88            | 47.39            |
| AVERAGE FADE - 17.66                                     |                  |                  |                  |
| SAMPLE D (CONTROL)                                       |                  |                  |                  |
| COLOUR   | 1 HOUR FADE D.E. | 2 HOUR FADE D.E. | 4 HOUR FADE D.E. |
| BLACK  | 4.03             | 6.31             | 17.45            |
| CYAN   | 27.92            | 40.03            | 66.43            |
| YELLOW   | 6.32             | 8.52             | 22.41            |
| MAGENTA  | 16.11            | 28.24            | 56.36            |
| AVERAGE FADE - 25.01                                     |                  |                  |                  |

Results show the lanthanum nitrate coated sheets exhibit less fade than the control (sample A) treated with nothing at all. The amine coated support (sample D) shows that the paper actually gets duller faster than no coating (sample A).

#### Example 5

Ink receiving sheets A and B were prepared in a similar way as described in Example 1. The coated layers on a polyester support had the following compositions:

|   | A                      | B                      |
|---|------------------------|------------------------|
| Gelatin with high is electric point                     | 4.200 g/m <sup>2</sup> | 4.200 g/m <sup>2</sup> |
| Latex LL 920/1  | 1.800 g/m <sup>2</sup> | 1.800 g/m <sup>2</sup> |
| Lanthanum nitrate                                       | 0.204 g/m <sup>2</sup> |                        |
| Olin 10G (idem example 1)                               | 0.175 g/m <sup>2</sup> | 0.175 g/m <sup>2</sup> |
| Crosslinker (idem example 1)                            | 0.015 g/m <sup>2</sup> | 0.075 g/m <sup>2</sup> |
| Latex LL 920/1 is available from Wacker Chemie, Germany |                        |                        |

The sheets were jetted with Hewlett Packard 500 series ink jet colour printing inks. Light stability was determined as described above in the testing procedures. The obtained results are reported in Table 5.

Table 5

| Ink receiving sheet | Key | Light Stability Loss in Density %<br>after 20 kJ Atlas |    |    |    |
|---------------------|-----|--|----|----|----|
|                     |     | Y  | M  | C  | K  |
| A                   | 1   | 17   | 24 | 26 | 3  |
| B                   | 2   | 20   | 46 | 64 | 13 |

Key:  
1 invention  
2 comparison

From these figures it can be clearly seen that the inks printed on the receiving sheet of the invention A fade much less than on the control receiving sheet B.

#### Example 6

Ink receiving sheets A and B were prepared in a similar way as described in Example 1. The coated layers on a polyester support had the following compositions:

|   | A                      | B                      |
|---|------------------------|------------------------|
| Hydroxyethyl cellulose (idem example 1) | 6.000 g/m <sup>2</sup> | 6.000 g/m <sup>2</sup> |
| Lanthanum nitrate                       | 0.500 g/m <sup>2</sup> |                        |
| Olin 10G (idem example 1)               | 0.210 g/m <sup>2</sup> | 0.210 g/m <sup>2</sup> |

The sheets were jetted with Hewlett Packard 500 series ink jet colour printing inks. Light stability was determined as described above in the testing procedures. The obtained results are reported in Table 6.

Table 6

| Ink receiving sheet | Key | Light Stability Loss in Density %<br>after 20 kJ Atlas |    |    |   |
|---------------------|-----|--|----|----|---|
|                     |     | Y  | M  | C  | K |
| A                   | 1   | 21   | 62 | 62 | 2 |
| B                   | 2   | 22   | 70 | 87 | 5 |

Key:

1 invention

2 comparison

From these figures it can be clearly seen that the inks printed on the receiving sheet A of the invention fade less than on the control receiving sheet B.

Example 7

Double layer ink receiving sheets A and B were prepared. The first layer was prepared in a similar way as described in Example 1. The first coated layer nearest to a RC-paper support had the following composition:

|                                     | A                      | B                      |
|-------------------------------------|------------------------|------------------------|
| Gelatin with high isoelectric point | 3.750 g/m <sup>2</sup> | 3.750 g/m <sup>2</sup> |
| Lanthanum nitrate                   | 0.300 g/m <sup>2</sup> |                        |
| Olin 10G (idem example 1)           | 0.125 g/m <sup>2</sup> | 0.125 g/m <sup>2</sup> |

The second layer was coated on this first layer, it had the following composition (same for A and B):

|                                     | A and B                |
|-------------------------------------|------------------------|
| Gelatin with high isoelectric point | 3.750 g/m <sup>2</sup> |
| Spacing agent                       | 0.100 g/m <sup>2</sup> |
| Olin 10G (idem example 1)           | 0.150 g/m <sup>2</sup> |
| Crosslinker (idem example 1)        | 0.033 g/m <sup>2</sup> |

Polystyrene beads of mean diameter 5 µm were used as spacing agent.

The sheets were jetted with Hewlett-Packard 500 series ink jet colour printing inks. Light stability was determined as described above in the testing procedures. The obtained results are reported in Table 7.

Table 7

| Ink receiving sheet | Key | Waterfastness Loss in Density %<br>after 1 min in water |    |   |   |
|---------------------|-----|---|----|---|---|
|                     |     | Y   | M  | C | K |
| A                   | 1   | 0   | 13 | 0 | 0 |
| B                   | 2   | 11  | 13 | 0 | 6 |

Key:  
1 invention  
2 comparison

From these figures it can be clearly seen that dye bleeding of the yellow and the black dye is much reduced when printed on the receiving sheet A of the invention compared to the control receiving sheet B.

Advantageously, the present invention provides image receiving media for use in ink jet recording which have excellent water resistance and lightfastness.

It will be recognised by those skilled in the art that the invention has wide application as a media which allows the possibility to print images intended to match those on silver halide photographic material. Further advantage is obtained by providing a recording medium which is suitable for office desk top publishing of colour graphics for ink jet printers.

The names 'X-rite', 'Tylose', 'Olin', 'Iris', 'Doothan', 'Lock Haven' and 'Melinex' are Registered Trade Marks.

In US 4146792, US 5011816 and US 5118349 there are described the use of certain rare earth metal complexes for their ability to fluoresce.

In G.B. 2147003 there is described a method of printing which comprises ink jet printing an image onto a medium comprising a support material containing in or on the surface portion thereof a water-soluble salt of a metal having a valency of from 2 to 4, together with a cationic organic material. Examples of water-soluble metal salts for use in G.B. 2147003 are salts of divalent calcium, divalent zinc, trivalent indium, trivalent aluminium cation, divalent magnesium and quadrivalent tin.

### Claims

1. An ink jet recording sheet on to which has been recorded an image by an ink jet printing device said recording sheet comprising a support having coated onto said support one or more layers receptive for aqueous inks, said recording sheet being characterised in that the coating comprises at least one water-soluble salt of Yttrium, Lanthanum, Cerium, Neodymium or Ytterbium.
2. An ink jet recording sheet according to claim 1 characterised in that the water-soluble salts of said metals are salts of mineral acids.
3. An ink jet recording sheet according to claim 2 characterised in that the water-soluble salts of said metals are salts of nitric acid.
4. An ink jet recording sheet according to claim 1 characterised in that lanthanum nitrate is present in the coating.
5. An ink jet recording sheet according to claim 1 characterised in that the water-soluble salts of said metals are salts of organic acids.
6. An ink jet recording sheet on to which has been recorded an image by an ink jet printing device said recording sheet comprising a support having coated onto said support one or more layers receptive for aqueous inks, said recording sheet being characterised in that its coating comprises at least one film forming hydrophilic polymer or a mixture of film forming hydrophilic polymers and imbedded in this film at least one water-soluble salt of Yttrium, Lanthanum, Cerium, Neodymium or Ytterbium.
7. An ink jet recording sheet according to claim 6 characterised in that the hydrophilic film forming polymer is selected from the group comprising natural polymers or modified products selected from albumin, gelatine, casein, starch, gum arabic, sodium alginate, hydroxyethyl cellulose, carboxymethyl cellulose;  $\alpha$ -,  $\beta$ -, or  $\gamma$ -cyclodextrine, polyvinyl alcohol; complete or partial saponified products of copolymers of vinyl acetate and other monomers; homopoly-



mers or copolymers with monomers of unsaturated carboxylic acids such as (meth)acrylic acid, maleic acid, crotonic acid and the like; homopolymers or copolymers with other vinyl monomers of sulfonated vinyl monomers such as vinylsulfonic acid, sulfonated styrene and the like; polyamides like polyvinylacrylamide, homopolymers or copolymers with other vinyl monomers of (meth)acrylamide, watersoluble nylon-type polymers; homopolymers or copolymers with monomers of ethylene oxide; polyethyleneimine; polyvinylpyrrolidone; polyurethane; polyester or mixtures of these compounds.

8. An ink jet recording sheet according to claim 6 characterised in that the hydrophilic film forming polymer is selected from the group comprising gelatine, casein, starch, gum arabic, sodium alginate, hydroxyethyl cellulose, carboxymethyl cellulose;  $\alpha$ -,  $\beta$ - or  $\gamma$ -cyclodextrine, polyvinyl alcohol; complete or partial saponified products of copolymers of vinyl acetate and other monomers; homopolymers or copolymers with monomers of unsaturated carboxylic acids such as (meth)acrylic acid, of sulfonated vinyl monomers such as vinylsulfonic acid, sulfonated styrene; homopolymers or copolymers with acrylamide, watersoluble nylon-type polymer, polyvinylpyrrolidone, polyurethane or mixtures of these compounds.
9. An ink jet recording sheet according to claim 6 characterised in that the hydrophilic film forming polymer is selected from the group comprising gelatine, starch, hydroxyethyl cellulose,  $\alpha$ -,  $\beta$ - or  $\gamma$ -cyclodextrine, polyvinyl alcohol; copolymers of vinyl acetate and other monomers; acrylamide, watersoluble nylon-type polymer, polyvinylpyrrolidone, polyurethane or mixtures of these compounds.
10. An ink jet recording sheet according to claim 1 characterised in that there is an ink-receiving layer on both sides of the support.
11. An ink jet recording sheet according to claim 10 characterised in that said layers have different compositions.
12. An ink jet recording sheet according to claim 6 characterised in that layers have a thickness of from 0.5 to 25 $\mu$ m.
13. An ink jet recording sheet according to claim 6 characterised in that the hydrophilic polymer in the layer or layers is crosslinked.
14. An ink jet recording sheet according to claim 13 characterised in that the crosslinking agents are selected from the group of formaldehyde, glyoxal, dihydroxydioxane, dichloro-hydroxy triazine, chloro- dihydroxy-triazine, 2-(4-dimethylcarbamoyl -pyridino) -ethane-sulfonate or 2,2'-Bis(vinylsulfonyl) -diethyl ether.
15. An ink jet recording sheet according to claim 6 characterised in that the amount of said metal salt is present in the range of 0.05 to 3.0g/m<sup>2</sup> in the hydrophilic polymer layer.
16. An ink jet recording sheet according to claim 15 characterised in that the amount of said metal salt present in the range of 0.1 to 0.9g/m<sup>2</sup>.
17. An ink jet recording sheet according to claim 6 characterised in that said hydrophilic polymer film contains in addition to said metal salts one or more non water soluble fillers or pigments.
18. An ink jet recording sheet according to claim 18 characterised in that the fillers or pigments are selected from clay, talc, zeolites, calcium-, barium- or magnesium-carbonate, calcium- or barium-sulphate, satin white, silicium oxide or colloidal silicium oxide.
19. An ink jet recording sheet according to claim 17 characterised in that the fillers or pigments are selected from satin white, silicium oxide or colloidal silicium oxide.
20. An ink jet recording sheet according to claim 19 characterised in that the filler is a non water soluble organic polymer.
21. An ink jet recording sheet according to claim 7 characterised in that said gelatine is acid pigskin gelatine, limed bone gelatine, acid or base hydrolysed gelatine or derivatised gelatine.
22. An ink jet recording sheet according to claim 21 characterised in that the derivatised gelatin is phthalated gelatine, carbamoylated gelatine, acetylated gelatine or trimellitic-acid modified gelatine.

23. An ink jet recording sheet according to claim 7 characterised in that the gelatine is a gelatine with an isoelectric point of 7 to 9.5.
- 5 24. An ink jet recording sheet according to claim 13 characterised in that said cross-linked gelatine film contains water-soluble salts of Yttrium, Lanthanum, Cerium, Neodymium or Ytterbium and a filler selected from clay, silicium oxide, colloidal silicium oxide, satin white or an organic polymer.
- 10 25. An ink jet recording sheet according to claim 13 characterised in that said gelatine film contains said water-soluble salts of Yttrium, Lanthanum, Cerium, Neodymium or Ytterbium and a cross-linked water soluble polymer or a mixture of cross-linked water soluble polymers selected from starch, hydroxyethyl cellulose,  $\alpha$ -,  $\beta$ - or  $\gamma$ -cyclodextrine, polyvinyl alcohol, vinyl acetate, acrylamide, water-soluble nylon-type polymer or polyvinylpyrrolidone.
26. An ink jet recording sheet according to claim 1 characterised in that said support is a clear thermoplastic film.
- 15 27. An ink jet recording sheet according to claim 1 characterised in that said support is an opaque thermoplastic film.
28. An ink jet recording sheet according to claim 1 characterised in that said support is a resin coated paper.
29. An ink jet recording sheet according to claim 1 characterised in that said support is plain paper.
- 20 30. An ink jet recording sheet according to claim 1 characterised in that said support is a surface treated plain paper.
31. An ink jet recording sheet according to claim 1 characterised in that said support is plain paper coated with a silica and polyvinyl alcohol matrix and which has been coated with an aqueous solution of at least one of said metal salts.
- 25 32. An ink jet recording sheet according to claim 30 characterised in that said aqueous solution is a 3-5% by weight solution of lanthanum nitrate.

#### Patentansprüche

- 30 1. Aufzeichnungsblatt für den Tintenstrahldruck, auf dem mit einer Tintenstrahldruckvorrichtung ein Bild aufgezeichnet worden ist, umfassend einen Träger mit einer oder mehreren darauf aufgebracht, für wäßrige Tinten aufnahmefähigen Schichten und gekennzeichnet dadurch, daß der Überzug mindestens ein wasserlösliches Yttrium-, Lanthan-, Cer-, Neodym- oder Ytterbiumsalz enthält.
- 35 2. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß es sich bei den wasserlöslichen Metallsalzen um Mineralsäuresalze handelt.
- 40 3. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 2, dadurch gekennzeichnet, daß es sich bei den wasserlöslichen Metallsalzen um Salpetersäuresalze handelt.
4. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß der Überzug Lanthannitrat enthält.
- 45 5. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß es sich bei den wasserlöslichen Metallsalzen um Salze von organischen Säuren handelt.
- 50 6. Aufzeichnungsblatt für den Tintenstrahldruck, auf dem mit einer Tintenstrahldruckvorrichtung ein Bild aufgezeichnet worden ist, umfassend einen Träger mit einer oder mehreren darauf aufgebracht, für wäßrige Tinten aufnahmefähigen Schichten und gekennzeichnet dadurch, daß der Überzug mindestens ein filmbildendes hydrophiles Polymer oder ein Gemisch von filmbildenden hydrophilen Polymeren enthält und in diesem Film mindestens ein wasserlösliches Yttrium-, Lanthan-, Cer-, Neodym- oder Ytterbiumsalz eingebettet ist.
- 55 7. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 6, dadurch gekennzeichnet, daß das hydrophile filmbildende Polymer aus der Gruppe bestehend aus natürlichen Polymeren oder modifizierten Produkten, ausgewählt unter Albumin, Gelatine, Casein, Stärke, Gummi arabicum, Natriumalginat, Hydroxyethylcellulose, Carboxymethylcellulose;  $\alpha$ -,  $\beta$ - oder  $\gamma$ -Cyclodextrin, Polyvinylalkohol; teil- oder vollverseiften Produkten von Copolymeren aus Vinylacetat und anderen Monomeren; Homopolymeren oder Copolymeren mit Monomeren ungesättigter Carbon-

säuren wie (Meth)acrylsäure, Maleinsäure, Crotonsäure und dergleichen; Homopolymeren von sulfonierten Vinylmonomeren wie Vinylsulfonsäure, sulfoniertem Styrol und dergleichen oder Copolymeren davon mit anderen Vinylmonomeren; Polyamiden wie Polyvinylacrylamid, Homopolymeren von (Meth)acrylamid oder Copolymeren davon mit anderen Vinylmonomeren, wasserlöslichen Polymeren vom Nylon-Typ; Homopolymeren oder Copolymeren mit Ethylenoxid-Monomeren; Polyethylenimin; Polyvinylpyrrolidon; Polyurethan; Polyester oder Gemischen dieser Verbindungen ausgewählt ist.

8. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 6, dadurch gekennzeichnet, daß das hydrophile filmbildende Polymer aus der Gruppe bestehend aus Gelatine, Casein, Stärke, Gummi arabicum, Natriumalginat, Hydroxyethylcellulose, Carboxymethylcellulose;  $\alpha$ -,  $\beta$ - oder  $\gamma$ -Cyclodextrin, Polyvinylalkohol; teil- oder vollverseiften Produkten von Copolymeren aus Vinylacetat und anderen Monomeren; Homopolymeren oder Copolymeren mit Monomeren ungesättigter Carbonsäuren wie (Meth)acrylsäure, von sulfonierten Vinylmonomeren wie Vinylsulfonsäure, sulfoniertem Styrol; Homopolymeren oder Copolymeren mit Acrylamid, wasserlöslichen Polymeren vom Nylon-Typ, Polyvinylpyrrolidon, Polyurethan oder Gemischen dieser Verbindungen ausgewählt ist.
9. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 6, dadurch gekennzeichnet, daß das hydrophile filmbildende Polymer aus der Gruppe bestehend aus Gelatine, Stärke, Hydroxyethylcellulose,  $\alpha$ -,  $\beta$ - oder  $\gamma$ -Cyclodextrin, Polyvinylalkohol; Copolymeren aus Vinylacetat und anderen Monomeren; Acrylamid, wasserlöslichen Polymeren vom Nylon-Typ, Polyvinylpyrrolidon, Polyurethan oder Gemischen dieser Verbindungen ausgewählt ist.
10. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß sich auf beiden Seiten des Trägers eine tintenempfangende Schicht befindet.
11. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 10, dadurch gekennzeichnet, daß die Schichten unterschiedliche Zusammensetzungen haben.
12. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 6, dadurch gekennzeichnet, daß die Schichten 0,5 bis 25  $\mu$ m dick sind.
13. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 6, dadurch gekennzeichnet, daß das hydrophile Polymer in der Schicht bzw. den Schichten vernetzt ist.
14. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 13, dadurch gekennzeichnet, daß die Vernetzer aus der Gruppe bestehend aus Formaldehyd, Glyoxal, Dihydroxydioxan, Dichlorhydroxytriazin, Chlordihydroxytriazin, 2-(4-Dimethylcarbamoylpyridino)ethansulfonat oder 2,2'-Bis(vinylsulfonyl)diethylether ausgewählt sind.
15. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 6, dadurch gekennzeichnet, daß das Metallsalz in der Schicht aus hydrophilem Polymer in einer Menge im Bereich von 0,05 bis 3,0 g/m<sup>2</sup> vorliegt.
16. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 15, dadurch gekennzeichnet, daß das Metallsalz in einer Menge im Bereich von 0,1 bis 0,9 g/m<sup>2</sup> vorliegt.
17. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 6, dadurch gekennzeichnet, daß der Film aus hydrophilem Polymer neben den Metallsalzen ein oder mehrere nicht wasserlösliche Füllstoffe oder Pigmente enthält.
18. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 17, dadurch gekennzeichnet, daß die Füllstoffe oder Pigmente unter Ton, Talk, Zeolithen, Calcium-, Barium- oder Magnesiumcarbonat, Calcium- oder Bariumsulfat, Satinweiß, Siliciumoxid oder kolloidalem Siliciumoxid ausgewählt sind.
19. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 17, dadurch gekennzeichnet, daß die Füllstoffe oder Pigmente unter Satinweiß, Siliciumoxid oder kolloidalem Siliciumoxid ausgewählt sind.
20. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 19, dadurch gekennzeichnet, daß es sich bei dem Füllstoff um ein nicht wasserlösliches organisches Polymer handelt.
21. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 7, dadurch gekennzeichnet, daß es sich bei der Gelatine um durch sauren Aufschluß von Schweinehaut gewonnene Gelatine, durch Äschern von Knochen gewonnene Gelatine, sauer oder basisch hydrolysierte Gelatine oder derivatisierte Gelatine handelt.

22. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 21, dadurch gekennzeichnet, daß es sich bei der derivatisierten Gelatine um phthalierte Gelatine, carbamoylierte Gelatine, acetylierte Gelatine oder mit Trimellitsäure modifizierte Gelatine handelt.
- 5 23. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 7, dadurch gekennzeichnet, daß es sich bei der Gelatine um eine Gelatine mit einem isoelektrischen Punkt von 7 bis 9,5 handelt.
24. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 13, dadurch gekennzeichnet, daß der Film aus vernetzter Gelatine wasserlösliche Yttrium-, Lanthan-, Cer-, Neodym- oder Ytterbiumsalze und einen Füllstoff, ausgewählt unter Ton, Siliciumoxid, kolloidalem Siliciumoxid, Satinweiß oder einem organischen Polymer, enthält.
- 10 25. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 13, dadurch gekennzeichnet, daß der Film aus Gelatine die wasserlöslichen Yttrium-, Lanthan-, Cer-, Neodym- oder Ytterbiumsalze und ein vernetztes wasserlösliches Polymer oder ein Gemisch aus vernetzten wasserlöslichen Polymeren, ausgewählt unter Stärke, Hydroxyethylcellulose,  $\alpha$ -,  $\beta$ - oder  $\gamma$ -Cyclodextrin, Polyvinylalkohol, Vinylacetat, Acrylamid, wasserlöslichen Polymeren vom Nylon-Typ oder Polyvinylpyrrolidon, enthält.
- 15 26. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß es sich bei dem Träger um eine klare thermoplastische Folie handelt.
- 20 27. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß es sich bei dem Träger um eine opake thermoplastische Folie handelt.
- 25 28. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß es sich bei dem Träger um ein harzbeschichtetes Papier handelt.
29. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß es sich bei dem Träger um Normalpapier handelt.
- 30 30. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß es sich bei dem Träger um oberflächenbehandeltes Normalpapier handelt.
31. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 1, dadurch gekennzeichnet, daß es sich bei dem Träger um mit einer Siliciumdioxid- und Polyvinylalkoholmatrix überzogenes Normalpapier handelt, das mit einer wäßrigen Lösung mindestens eines der Metallsalze überzogen worden ist.
- 35 32. Aufzeichnungsblatt für den Tintenstrahldruck nach Anspruch 30, dadurch gekennzeichnet, daß es sich bei der wäßrigen Lösung um eine 3-5 gew.-%ige Lösung von Lanthan-nitrat handelt.

#### 40 Revendications

1. Feuille d'enregistrement de jet d'encre sur laquelle une image a été enregistrée par un dispositif d'impression à jet d'encre, ladite feuille d'enregistrement comprenant un support, ledit support étant revêtu d'une ou de plusieurs couches sensibles aux encres aqueuses, ladite feuille d'enregistrement étant caractérisée en ce que le revêtement comprend au moins un sel soluble dans l'eau d'yttrium, de lanthane, de cérium, de néodyme ou d'ytterbium.
- 45 2. Feuille d'enregistrement de jet d'encre suivant la revendication 1, caractérisée en ce que les sels solubles dans l'eau desdits métaux sont des sels d'acides minéraux.
- 50 3. Feuille d'enregistrement de jet d'encre suivant la revendication 2, caractérisée en ce que les sels solubles dans l'eau desdits métaux sont des sels d'acide nitrique.
4. Feuille d'enregistrement de jet d'encre suivant la revendication 1, caractérisée en ce que le nitrate de lanthane est présent dans le revêtement.
- 55 5. Feuille d'enregistrement de jet d'encre suivant la revendication 1, caractérisée en ce que les sels solubles dans l'eau desdits métaux sont des sels d'acides organiques.

6. Feuille d'enregistrement de jet d'encre sur laquelle une image a été enregistrée par un dispositif d'impression à jet d'encre, ladite feuille d'enregistrement comprenant un support, ledit support étant revêtu d'une ou de plusieurs couches sensibles aux encres aqueuses, ladite feuille d'enregistrement étant caractérisée en ce que son revêtement comprend au moins un polymère hydrophile filmogène ou un mélange de polymères hydrophiles filmogènes et, noyé dans ce film, au moins un sel soluble dans l'eau d'yttrium, de lanthane, de cérium, de néodyme ou d'ytterbium.
7. Feuille d'enregistrement de jet d'encre suivant la revendication 6, caractérisée en ce que le polymère filmogène hydrophile est choisi dans le groupe comprenant des polymères naturels ou des produits modifiés choisis parmi l'albumine, la gélatine, la caséine, l'amidon, la gomme arabique, l'alginate de sodium, l'hydroxyéthylcellulose, la carboxyméthylcellulose; l' $\alpha$ -,  $\beta$ -, ou  $\gamma$ -cyclodextrine, l'alcool polyvinylique; des produits partiellement ou entièrement saponifiés de copolymères d'acétate de vinyle et d'autres monomères; des homopolymères ou des copolymères avec des monomères d'acides carboxyliques insaturés tels que l'acide (méth)acrylique, l'acide maléique, l'acide crotonique et autres; des homopolymères de copolymères avec d'autres monomères de vinyle de monomères de vinyle sulfonés tels que l'acide vinylsulfonique, le styrène sulfoné et autres; des polyamides comme l'acrylamide polyvinylique, des homopolymères ou des copolymères avec d'autres monomères de vinyle de (méth)acrylamide, des polymères de type Nylon solubles dans l'eau; des homopolymères ou des copolymères avec des monomères d'oxyde d'éthylène; le polyéthylèneimine; la polyvinylpyrrolidone; le polyuréthane; le polyester ou des mélanges de ces composés.
8. Feuille d'enregistrement de jet d'encre suivant la revendication 6, caractérisée en ce que le polymère hydrophile filmogène est choisi dans le groupe comprenant la gélatine, la caséine, l'amidon, la gomme arabique, l'alginate de sodium, l'hydroxyéthylcellulose, la carboxyméthylcellulose; l' $\alpha$ -,  $\beta$ -, ou  $\gamma$ -cyclodextrine, l'alcool polyvinylique; des produits partiellement ou entièrement saponifiés de copolymères d'acétate de vinyle et d'autres monomères; des homopolymères ou des copolymères avec des monomères d'acides carboxyliques insaturés tels que l'acide (méth)acrylique, de monomères de vinyle sulfonés tels que l'acide vinylsulfonique, le styrène sulfoné; des homopolymères ou des copolymères avec de l'acrylamide, un polymère de type Nylon soluble dans l'eau, la polyvinylpyrrolidone, le polyuréthane ou des mélanges de ces composés.
9. Feuille d'enregistrement de jet d'encre suivant la revendication 6, caractérisée en ce que le polymère hydrophile filmogène est choisi dans le groupe comprenant la gélatine, l'amidon, l'hydroxyéthylcellulose, l' $\alpha$ -,  $\beta$ -, ou  $\gamma$ -cyclodextrine, l'alcool polyvinylique; des copolymères d'acétate de vinyle et d'autres monomères; l'acrylamide, un polymère de type Nylon soluble dans l'eau, la polyvinylpyrrolidone, le polyuréthane ou des mélanges de ces composés.
10. Feuille d'enregistrement de jet d'encre suivant la revendication 1, caractérisée en ce qu'une couche recevant de l'encre est prévue sur les deux côtés du support.
11. Feuille d'enregistrement de jet d'encre suivant la revendication 10, caractérisée en ce que lesdites couches présentent différentes compositions.
12. Feuille d'enregistrement de jet d'encre suivant la revendication 6, caractérisée en ce que les couches présentent une épaisseur comprise entre 0,5 et 25  $\mu\text{m}$ .
13. Feuille d'enregistrement de jet d'encre suivant la revendication 6, caractérisée en ce que le polymère hydrophile dans la ou les couches est réticulé.
14. Feuille d'enregistrement de jet d'encre suivant la revendication 13, caractérisée en ce que les agents de réticulation sont choisis dans le groupe comprenant le formaldéhyde, le glyoxal, le dihydroxydioxane, la dichlorohydroxytriazine, la chlorodihydroxytriazine, le sulfonate de 2-(4-diméthylcarbamoylepyridino)éthane ou l'éther de 2,2'-bis(vinyl-sulfonyl)diéthyle.
15. Feuille d'enregistrement de jet d'encre suivant la revendication 6, caractérisée en ce que ledit sel métallique est présent en quantité comprise entre 0,05 et 3,0 g/m<sup>2</sup> dans la couche de polymère hydrophile.
16. Feuille d'enregistrement de jet d'encre suivant la revendication 15, caractérisée en ce que ledit sel métallique est présent en quantité comprise entre 0,1 et 0,9 g/m<sup>2</sup>.

17. Feuille d'enregistrement de jet d'encre suivant la revendication 6, caractérisée en ce que ledit film de polymère hydrophile contient, outre lesdits sels métalliques, une ou plusieurs matières de charge ou pigments non solubles dans l'eau.
- 5 18. Feuille d'enregistrement de jet d'encre suivant la revendication 18, caractérisée en ce que les matières de charge ou pigments sont choisis parmi l'argile, le talc, les zéolites, le carbonate de calcium, de baryum ou de magnésium, le sulfate de calcium ou de baryum, le blanc satiné, l'oxyde de silicium ou l'oxyde de silicium colloïdal.
- 10 19. Feuille d'enregistrement de jet d'encre suivant la revendication 17, caractérisée en ce que les matières de charge ou pigments sont choisis parmi le blanc satiné, l'oxyde de silicium ou l'oxyde de silicium colloïdal.
20. Feuille d'enregistrement de jet d'encre suivant la revendication 19, caractérisée en ce que la matière de charge est un polymère organique non soluble dans l'eau.
- 15 21. Feuille d'enregistrement de jet d'encre suivant la revendication 7, caractérisée en ce que ladite gélatine est de la gélatine de derme de porc acide, de la gélatine d'os chaulée, de la gélatine hydrolysée acide ou base ou de la gélatine obtenue par dérivatisation.
- 20 22. Feuille d'enregistrement de jet d'encre suivant la revendication 21, caractérisée en ce que la gélatine obtenue par dérivatisation est de la gélatine à base de phtalate, de la gélatine de carbamoylate, de la gélatine acétylée ou de la gélatine modifiée à l'acide trimellitique.
- 25 23. Feuille d'enregistrement de jet d'encre suivant la revendication 7, caractérisée en ce que la gélatine est une gélatine comportant un point isoélectrique compris entre 7 et 9,5.
- 30 24. Feuille d'enregistrement suivant la revendication 13, caractérisée en ce que ledit film de gélatine réticulé contient des sels solubles dans l'eau d'yttrium, de lanthane, de cérium, de néodyme ou d'ytterbium et une matière de charge choisie parmi l'argile, l'oxyde de silicium, l'oxyde de silicium colloïdal, le blanc satiné ou un polymère organique.
- 35 25. Feuille d'enregistrement de jet d'encre suivant la revendication 13, caractérisée en ce que ledit film de gélatine contient lesdits sels solubles dans l'eau d'yttrium, de lanthane, de cérium, de néodyme ou d'ytterbium et un polymère soluble dans l'eau réticulé ou un mélange de polymères solubles dans l'eau réticulés choisis parmi l'amidon, l'hydroxyéthylcellulose, l' $\alpha$ -,  $\beta$ -, ou  $\gamma$ -cyclodextrine, l'alcool polyvinylique, l'acétate de vinyle, l'acrylamide, un polymère de type Nylon soluble dans l'eau ou la polyvinylpyrrolidone.
- 40 26. Feuille d'enregistrement de jet d'encre suivant la revendication 1, caractérisée en ce que ledit support est un film thermoplastique transparent.
- 45 27. Feuille d'enregistrement pour jet d'encre selon la revendication 1, caractérisée en ce que ledit support est un film thermoplastique opaque.
28. Feuille d'enregistrement pour jet d'encre selon la revendication 1, caractérisée en ce que ledit support est du papier enduit de résine.
- 50 29. Feuille d'enregistrement pour jet d'encre selon la revendication 1, caractérisée en ce que ledit support est du papier uni.
30. Feuille d'enregistrement pour jet d'encre selon la revendication 1, caractérisée en ce que ledit support est du papier uni traité en surface.
- 55 31. Feuille d'enregistrement pour jet d'encre selon la revendication 1, caractérisée en ce que ledit support est du papier uni enduit d'une matrice de silice et d'alcool polyvinylique et qui a été enduit d'une solution aqueuse d'au moins l'un desdits sels métalliques.
32. Feuille d'enregistrement pour jet d'encre selon la revendication 30, caractérisée en ce que ladite solution aqueuse est une solution à 3-5 % en poids de nitrate de lanthane.